

10/523,472
Updated Search
LYCOOK 8/6/07

d his

(FILE 'HOME' ENTERED AT 13:48:12 ON 06 AUG 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 13:48:31 ON 06
AUG 2007

L1 47780 S (HEPATITIS B SURFACE)
L2 17 S L1 AND (ALUMINIUM HYDROXIDE)
L3 10 DUPLICATE REMOVE L2 (7 DUPLICATES REMOVED)

=>

ANSWER 5 OF 10 BIOSIS COPYRIGHT (c) 2007 The Thomson Corporation on STN

DUPLICATE 4

AN 1999:63533 BIOSIS

DN PREV199900063533

TI Evidence for the denaturation of recombinant hepatitis B surface antigen on aluminium hydroxide gel.

AU Tleugabulova, Dina [Reprint author]; Falcon, Viviana; Penton, Eduardo

CS Quality Control Dep., Natl. Cent. Bioprod., P.O. Box 6048, Havana 6, Cuba

SO Journal of Chromatography B, (Dec. 11, 1998) Vol. 720, No. 1-2, pp.

153-163. print.

CODEN: JCBADL. ISSN: 0378-4347.

DT Article

LA English

ED Entered STN: 16 Feb 1999

Last Updated on STN: 16 Feb 1999

AB Despite the complexity of the subject of protein-alum interactions, a valuable information can be obtained by analyzing the adsorbed and desorbed protein by common physico-chemical methods. In the present work, to approach the adsorption of hepatitis B surface antigen (HBsAg) on alum, the experimental data were supported by complementary analyses of the adsorbed protein by immunoelectron microscopy and the desorbed protein by denaturing size-exclusion chromatography and sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions. First, the depletion of HBsAg was investigated. The aspects assessed were the conditions, recovery and chromatographic performance of the desorbed protein. The results obtained strongly suggested the loss of particulate structure of HBsAg after adsorption on alum. This conclusion was further reinforced by direct immunoelectron microscopic visualization of HBsAg in the adsorbed state.

CC Pharmacology - Immunological processes and allergy 22018

Comparative biochemistry 10010

Biochemistry methods - Proteins, peptides and amino acids 10054

Biochemistry studies - General 10060

Biochemistry studies - Proteins, peptides and amino acids 10064

Biochemistry studies - Minerals 10069

Biophysics - Methods and techniques 10504

Biophysics - Molecular properties and macromolecules 10506

Pharmacology - Clinical pharmacology 22005

Virology - Animal host viruses 33506

Immunology - Bacterial, viral and fungal 34504

Medical and clinical microbiology - Virology 36006

IT Major Concepts

Biochemistry and Molecular Biophysics; Methods and Techniques;
Pharmaceuticals (Pharmacology)

IT Chemicals & Biochemicals

alum; aluminum hydroxide gels; proteins: analysis; recombinant hepatitis B surface antigen: analysis, denaturation; vaccines: analysis

IT Methods & Equipment

immunoelectron microscopy: analytical method, electron microscopy: CB, scanning electron microscopy; size exclusion chromatography: analytical method, liquid chromatography; SDS-polyacrylamide gel electrophoresis: analytical method, electrophoretic techniques, purification method; SDS-PAGE system: Hoefer Scientific Instrument, equipment

IT Miscellaneous Descriptors

protein-alum interactions: analysis

ORGN Classifier

Hepadnaviridae 03301

Super Taxa

DNA and RNA Reverse Transcribing Viruses; Viruses; Microorganisms

Organism Name

hepatitis B virus

Taxa Notes

DNA and RNA Reverse Transcribing Viruses, Microorganisms, Viruses

RN 10043-01-3Q (alum)
10043-67-1Q (alum)
21645-51-2 (ALUMINUM HYDROXIDE)

d his

(FILE 'HOME' ENTERED AT 13:48:12 ON 06 AUG 2007)

FILE 'BIOSIS, CAPLUS, EMBASE, MEDLINE, JAPIO' ENTERED AT 13:48:31 ON 06
AUG 2007

L1 47780 S (HEPATITIS B SURFACE)
L2 17 S L1 AND (ALUMINIUM HYDROXIDE)
L3 10 DUPLICATE REMOVE L2 (7 DUPLICATES REMOVED)

=>

DUPLICATE 4

AN 1999:63533 BIOSIS

DN PREV199900063533

TI Evidence for the denaturation of recombinant hepatitis B
surface antigen on aluminium hydroxide gel.

AU Tleugabulova, Dina [Reprint author]; Falcon, Viviana; Penton, Eduardo

CS Quality Control Dep., Natl. Cent. Bioprod., P.O. Box 6048, Havana 6, Cuba

SO Journal of Chromatography B, (Dec. 11, 1998) Vol. 720, No. 1-2, pp.
153-163. print.

CODEN: JCBADL. ISSN: 0378-4347.

DT Article

LA English

ED Entered STN: 16 Feb 1999

Last Updated on STN: 16 Feb 1999

AB Despite the complexity of the subject of protein-alum interactions, a valuable information can be obtained by analyzing the adsorbed and desorbed protein by common physico-chemical methods. In the present work, to approach the adsorption of hepatitis B surface antigen (HBsAg) on alum, the experimental data were supported by complementary analyses of the adsorbed protein by immunoelectron microscopy and the desorbed protein by denaturing size-exclusion chromatography and sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions. First, the depletion of HBsAg was investigated. The aspects assessed were the conditions, recovery and chromatographic performance of the desorbed protein. The results obtained strongly suggested the loss of particulate structure of HBsAg after adsorption on alum. This conclusion was further reinforced by direct immunoelectron microscopic visualization of HBsAg in the adsorbed state.

CC Pharmacology - Immunological processes and allergy 22018

Comparative biochemistry 10010

Biochemistry methods - Proteins, peptides and amino acids 10054

Biochemistry studies - General 10060

Biochemistry studies - Proteins, peptides and amino acids 10064

Biochemistry studies - Minerals 10069

Biophysics - Methods and techniques 10504

Biophysics - Molecular properties and macromolecules 10506

Pharmacology - Clinical pharmacology 22005

Virology - Animal host viruses 33506

Immunology - Bacterial, viral and fungal 34504

Medical and clinical microbiology - Virology 36006

IT Major Concepts

Biochemistry and Molecular Biophysics; Methods and Techniques;
Pharmaceuticals (Pharmacology)

IT Chemicals & Biochemicals

alum; aluminum hydroxide gels; proteins: analysis; recombinant
hepatitis B surface antigen: analysis,
denaturation; vaccines: analysis

IT Methods & Equipment

immunoelectron microscopy: analytical method, electron microscopy: CB,
scanning electron microscopy; size exclusion chromatography: analytical
method, liquid chromatography; SDS-polyacrylamide gel electrophoresis:
analytical method, electrophoretic techniques, purification method;
SDS-PAGE system: Hoefer Scientific Instrument, equipment

IT Miscellaneous Descriptors

protein-alum interactions: analysis

ORGN Classifier

Hepadnaviridae 03301

Super Taxa

DNA and RNA Reverse Transcribing Viruses; Viruses; Microorganisms

Organism Name

hepatitis B virus

Taxa Notes

DNA and RNA Reverse Transcribing Viruses, Microorganisms, Viruses

RN 10043-01-3Q (alum)
10043-67-1Q (alum)
21645-51-2 (ALUMINUM HYDROXIDE)

ANSWER 3 OF 10 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights reserved on STN DUPLICATE 2

AN 1999168176 EMBASE

TI Purification and characterization of hepatitis B virus surface antigen particles produced in Drosophila Schneider-2 cells.

AU Deml L.; Schirmbeck R.; Reimann J.; Wolf H.; Wagner R.

CS R. Wagner, Institute Medical Microbiology, Klinikum Regensburg, University of Regensburg, Franz-Josef-Strauss Allee 11, 95053 Regensburg, Germany. ralf.wagner@klinik.uni-regensburg.de

SO Journal of Virological Methods, (1999) Vol. 79, No. 2, pp. 205-217. . Refs: 52
ISSN: 0166-0934 CODEN: JVMEDH

PUI S 0166-0934(99)00022-1

CY Netherlands

DT Journal; Article

FS 026 Immunology, Serology and Transplantation
037 Drug Literature Index
004 Microbiology

LA English

SL English

ED Entered STN: 27 May 1999
Last Updated on STN: 27 May 1999

AB The small surface antigen of hepatitis B virus (HBV) was produced in Drosophila melanogaster Schneider-2 (DS-2) cells transfected stably using an inducible Drosophila metallothionein promoter. Selected clonal DS-2 cell-lines expressed and secreted large quantities of HBsAg particles consisting exclusively of non-glycosylated 25 kDa proteins. HBsAg produced by DS-2 cells had physical and biochemical properties very similar to 22 nm particles derived from the human hepatoma cell-line PLC/PRF/5. DS-2 cell-derived HBsAg particles were purified near homogeneity by a strategy based on protein concentration, precipitation and ultracentrifugation. The resulting HBsAg product was <98% pure. A single immunisation of BALB/c mice with both DS-2 and yeast-cell derived purified HBsAg particles without adjuvants elicited a substantial humoral antibody and class-I restricted cytotoxic T lymphocyte (CTL) response. Adsorption of HBsAg particles to aluminium hydroxide resulted in increased levels of HBsAg-specific antibodies. However, CTLs were not elicited by HBsAg/Alum combinations. Thus, stably transfected DS-2 cells provide a useful source for the production of HBV subviral particles for diagnostic and research purposes as well as for novel vaccine development. Copyright (C) 1999 Elsevier Science B.V.

CT Medical Descriptors:
*hepatitis b virus
*antigen expression
*immunogenicity
gene expression regulation
drosophila melanogaster
ultracentrifugation
antibody response
gene expression system
immunization
hepatitis b: PC, prevention
immunoblotting
antigenicity
cytotoxic t lymphocyte
nonhuman
mouse
animal experiment
controlled study
animal cell
article
priority journal
Drug Descriptors:
*hepatitis b surface antigen

*hepatitis b vaccine: DV, drug development

ANSWER 3 OF 10 EMBASE COPYRIGHT (c) 2007 Elsevier B.V. All rights reserved on STN DUPLICATE 2

AN 1999168176 EMBASE
TI Purification and characterization of hepatitis B virus surface antigen particles produced in Drosophila Schneider-2 cells.
AU Deml L.; Schirmbeck R.; Reimann J.; Wolf H.; Wagner R.
CS R. Wagner, Institute Medical Microbiology, Klinikum Regensburg, University of Regensburg, Franz-Josef-Strauss Allee 11, 95053 Regensburg, Germany. ralf.wagner@klinik.uni-regensburg.de
SO Journal of Virological Methods, (1999) Vol. 79, No. 2, pp. 205-217. .
Refs: 52
ISSN: 0166-0934 CODEN: JMVMDH
PUI S 0166-0934(99)00022-1
CY Netherlands
DT Journal; Article
FS 026 Immunology, Serology and Transplantation
037 Drug Literature Index
004 Microbiology
LA English
SL English
ED Entered STN: 27 May 1999
Last Updated on STN: 27 May 1999
AB The small surface antigen of hepatitis B virus (HBV) was produced in Drosophila melanogaster Schneider-2 (DS-2) cells transfected stably using an inducible Drosophila metallothionein promoter. Selected clonal DS-2 cell-lines expressed and secreted large quantities of HBsAg particles consisting exclusively of non-glycosylated 25 kDa proteins. HBsAg produced by DS-2 cells had physical and biochemical properties very similar to 22 nm particles derived from the human hepatoma cell-line PLC/PRF/5. DS-2 cell-derived HBsAg particles were purified near homogeneity by a strategy based on protein concentration, precipitation and ultracentrifugation. The resulting HBsAg product was <98% pure. A single immunisation of BALB/c mice with both DS-2 and yeast-cell derived purified HBsAg particles without adjuvants elicited a substantial humoral antibody and class-I restricted cytotoxic T lymphocyte (CTL) response. Adsorption of HBsAg particles to aluminium hydroxide resulted in increased levels of HBsAg-specific antibodies. However, CTLs were not elicited by HBsAg/Alum combinations. Thus, stably transfected DS-2 cells provide a useful source for the production of HBV subviral particles for diagnostic and research purposes as well as for novel vaccine development. Copyright (C) 1999 Elsevier Science B.V.
CT Medical Descriptors:
*hepatitis b virus
*antigen expression
*immunogenicity
gene expression regulation
drosophila melanogaster
ultracentrifugation
antibody response
gene expression system
immunization
hepatitis b: PC, prevention
immunoblotting
antigenicity
cytotoxic t lymphocyte
nonhuman
mouse
animal experiment
controlled study
animal cell
article
priority journal
Drug Descriptors:
*hepatitis b surface antigen

→ May 1999

*hepatitis b vaccine: DV, drug development